## **AMENDMENTS TO THE CLAIMS:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **LISTING OF CLAIMS:**

1. (Currently Amended) A moldable implant mass composition for use in repairing a bone defect in a living organism, comprising:

a plurality of biocompatible synthetic non-polymeric granules, said granules having an equivalent diameter of about 100 µm to about 4,000 µm;

implant mass comprising a composite matrix of the granules bound together by the biocompatible polymer, and macropores between adjacent granules, said granules so as to form an implant mass comprising a plurality of distinct granules coated with said biocompatible polymer, said biocompatible polymer comprising about 4% to about 20% of the total weight of the implant mass; and

a plasticizer in said implant mass in an amount sufficient to condition at least a portion of said biocompatible polymer so that said implant mass is initially plastically deformable into a desired shape and then hardenable upon removal of at least a portion of said plasticizer from said implant mass.

2. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, wherein the granules comprise at least one material selected from the group consisting of biocompatible ceramics, and biocompatible glasses.

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- 3. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, wherein the granules comprise a at least one material selected from the group consisting of silicon oxide, calcium sulphate, and calcium phosphate.
- 4. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, wherein the granules comprise a at least one material selected from the group consisting of monocalcium phosphate monohydrate, monocalcium phosphate anhydrous, dicalcium phosphate dihydrate, dicalcium phosphate anhydrous, tetracalcium phosphate, calcium orthophosphate phosphate, calcium pyrophosphate, α-tricalcium phosphate, β-tricalcium phosphate, hydroxyapatite, carbonate hydroxyapatite, apatite, and bioglass.
- (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, wherein the granules are biodegradable.
- 6. (Currently Amended) A moldable implant <u>mass</u> composition as in defined claim 1, wherein said biocompatible polymer is biodegradable.
- 7. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, wherein said biocompatible polymer is comprises at least one polymer selected from the group consisting of poly(α-hydroxyesters), poly(orthoesters), poly(ether esters), polyanhydrides, poly(phosphazenes), poly(propylene fumarates), poly(ester amides), poly(ethylene fumarates), poly(amino acids), polysaccharides, polypeptides, poly(hydroxy butyrates), poly(hydroxy valerates), polyurethanes,

poly(malic acid), polylactides, polyglycolides, poly(lactide-co-glycolide), polycaprolactones, poly(glycolide-co-trimethylene carbonates), and polydioxanones.

- 8. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, wherein the biocompatible polymer comprises poly(lactide-co-glycolide).
- 9. (Currently Amended) A moldable implant <u>mass</u> composition as in claim 1, wherein said plasticizer is selected from the group consisting of n-methyl-2-pyrrolidone, acetone, ethyl lactate, ethyl acetate, ethyl formiate, acetyltributylcitrate, triethyl citrate, lactic acid, citric acid tetrahydrofuran, toluene, alcohol and carbon dioxide.
- 10. (Currently Amended) A moldable implant <u>mass</u> composition as in defined in claim 1, further comprising a biologically active substance.
- 11. (Currently Amended) A moldable implant <u>mass</u> composition as in defined <u>in</u> claim 1, wherein said plasticizer is extractable from said implant mass when contacted with a hardener.
- 12. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 11, wherein said hardener comprises water or a body fluid.

13. - 14. (Canceled)

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- 15. (Previously Presented) The composite matrix of claim 43, further comprising a membrane on a surface of said composite matrix.
- 16. (Currently Amended) A moldable implant <u>mass</u> composition as defined in claim 1, in combination with a syringe that is capable of injecting the moldable implant composition into a bone defect.
  - 17. 40. (Canceled)
  - 41. (Currently Amended) A composite implant mass comprising:

a structural component, the structural component comprising a plurality of biocompatible synthetic non-polymeric granules, the granules being regularly-sized, regularly shaped, or spherical, and the granules having an equivalent diameter of about 100 µm to about 4,000 µm;

a biocompatible polymer on at least a portion of <u>each of</u> the granules; and a plasticizer in an amount sufficient to condition at least a portion of the biocompatible polymer so that the granules of the implant mass are bound together by the biocompatible polymer, and the implant mass is <u>initially</u> plastically deformable.

42. (Previously Presented) The implant mass of claim 41, wherein the biocompatible polymer comprises 4% to 20% of the total weight of the implant mass.

43. (Currently Amended) A composite matrix comprising:

a structural matrix, the structural matrix comprising a plurality of biocompatible synthetic non-polymeric granules bound together, at least in part, by a biocompatible polymer coating formed on each <u>of the</u> granules; and

an open porous region comprising macropores between adjacent coated granules;

wherein the structural matrix does not contain any bone particles.

- 44. (Previously Presented) The composite matrix of claim 43, wherein the open porous region is filled with air or gas.
- 45. (Previously Presented) The composite matrix of claim 43, wherein the open porous region is filled with a liquid, solid particles, or a gel.
- 46. (Previously Presented) The composite matrix of claim 43, wherein the biocompatible polymer comprises 4% to 20% of the total weight of the composite.
- 47. (Currently Amended) The moldable implant <u>mass</u> composition as defined in claim 1, wherein the granules are regularly-shaped, regularly-sized, or spherical.
- 48. (Currently Amended) The moldable implant <u>mass</u> composition as defined in claim 47, wherein the granules have an equivalent diameter of about 100

 $\mu m$  to about 4,000  $\mu m$  and the polymer coating has a thickness of about 1  $\mu m$  to about 300  $\mu m$ .

- 49. (Currently Amended) The moldable implant <u>mass</u> composition as defined in claim 47, wherein the granules have an equivalent diameter of about 500 μm to about 1,500 μm, and the polymer coating has a thickness of about 5 μm to about 30 μm.
- 50. (Currently Amended) The moldable implant <u>mass</u> composition as claimed in claim 1, wherein the implant composition in claim 1, wherein the implant composition does not contain bone particles.
- 51. (Previously Presented) The implant mass of claim 41, wherein the granules have an equivalent diameter of about 500 μm to about 1,500 μm.
- 52. (Previously Presented) The implant mass of claim 41, wherein the granules have a coating of the polymer and are distinct from one another.
- 53. (Previously Presented) The implant mass of claim 52, wherein the coating has a thickness of about 1  $\mu m$  to about 30  $\mu m$ .
- 54. (Previously Presented) The implant mass of claim 41, wherein the coating has a thickness of about 5 μm to about 30 μm.

- 55. (Previously Presented) The composite matrix of claim 43, wherein the granules are regularly-sized, regularly-shaped, or spherical.
- 56. (Currently Amended) The moldable implant <u>mass</u> composition as defined in claim 1, wherein the macropores have an average diameter of about greater than 10  $\mu$ m to about 2000  $\mu$ m.
- 57. (Currently Amended) The moldable implant <u>mass</u> composition as defined in claim 56, wherein the macropores have an average diameter of about 100 μm to about 500 μm.
- 58. (Currently Amended) The moldable implant <u>mass</u> composition is defined in claim 1, wherein the granules or biocompatible polymer comprise micropores.
- 59. (Currently Amended) The moldable implant <u>mass</u> composition as defined in claim 1, wherein the granules comprise calcium phosphate.
- 60. (Currently Amended) The moldable implant mass composition of claim
  59, wherein the calcium phosphate comprises β-tricalciumphosphate or
  hydroxyapatite.

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- 61. (Currently Amended) The moldable implant <u>mass</u> composition of claim 58, wherein the biocompatible polymer comprises polylactide-co-glycolide, and the plasticizer comprises n-methyl-2-pyrrolidone, acetone, or an alcohol.
- 62. (Currently Amended) The moldable implant mass <u>composition</u> of claim 1, wherein the granules comprise regularly-shaped spherical particles having a homogenous coating of the biocompatible polymer.
- 63. (Previously Presented) The composite matrix of claim 43, wherein the macropores have an average diameter of about greater than 10  $\mu$ m to about 2000  $\mu$ m.
- 64. (Previously Presented) The composite matrix of claim 63, wherein the macropores have an average diameter of about 100 μm to about 500 μm.